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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/073,138	02/13/2002	Darrell R. Anderson	P 0280705 7969 1995-30-0233CP1	
7590 10/06/2004		EXAMINER		
Pillsbury Winthrop LLP			GAMBEL, PHILLIP	
Intellectual Prop	perty Group			
1600 Tysons Boulevard			ART UNIT	PAPER NUMBER
McLean, VA 22102			1644	
			DATE MAILED: 10/06/2004	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
Office Action Summers	10/073,138	ANDERSON ET AL.				
Office Action Summary	Examiner	Art Unit				
	Phillip Gambel	1644				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE						
closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims Claim(s) \(\frac{18}{20} \) is/are pending in the application 4a) Of the above claim(s) is/are withdraw 5) \(\begin{aligned} \text{Claim(s)} & is/are allowed. \\ 6) \(\begin{aligned} \text{Claim(s)} & is/are rejected. \\ 7) \(\begin{aligned} \text{Claim(s)} & is/are objected to. \\ 8) \(\begin{aligned} \text{Claim(s)} & is/are subject to restriction and/or is/are subject to restriction and/or is/are subject to restriction and/or is/are subject to restriction. \]	n. vn from consideration.					
Application Papers		•				
9) The specification is objected to by the Examiner 10) The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the or Replacement drawing sheet(s) including the correction 11) The oath or declaration is objected to by the Examiner	epted or b) objected to by the Edrawing(s) be held in abeyance. See on is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).				
Priority under 35 U.S.C. § 119						
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 						
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal Pa					

Application/Control Number: 10/073,138

Art Unit: 1644

DETAILED ACTION

- 1. This application appears to be in compliance with the Sequence Rules.
- 2. Prior to setting forth the restriction requirement, it is pointed out that the claims are drawn to patentably distinct products and methods, which rely upon differ ingredients, process steps and endpoints, which, in turn, require non-coextensive searches to such an extent that they are considered separately patentable. Both the claimed products and methods rely upon antibodies that bind B7-1 or B7-2. B7-1 and B7-2 differ in structure, expression and modes of action to such an extent and require non-coextensive searches to such an extent that they are considered separately patentable. Therefore, the restriction will be set forth for each of the various Groups, irrespective of the format of the claims.
- 3. Restriction to one of the following inventions is required under 35 U.S.C. § 121:
- I. Claims 1-3, 6, 7, 9-14 and 26, drawn to anti-B7-1 antibodies and compositions thereof, classified in Class 424, subclass 153.1 and Class 530, subclass 387.1.
- II. Claims 1, 4-6, 8-13, 15 and 26, drawn to anti-B7-2 antibodies and compositions thereof, classified in Class 424, subclass 141.1 and Class 530, subclass 388.1.
- III. Claims 16, 18, 20, 22, 24 and, 27, drawn to a method of treating a disease with anti-B7-1 antibodies, classified in Class 424, subclass 130.1.
- IV. Claims 17, 19, 21, 23, 25 and 28, drawn to a method of treating a disease with anti-B7-2 antibodies, classified in Class 424, subclass 173.1.
- 4. (Inventions I and III) and (Inventions II and IV) are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (M.P.E.P. § 806.05(h)).

In the instant case, the products as claimed can be used in a materially different process such as affinity purification or detection assays .

- 5. Inventions I and II are different methods, which require different ingredients, process steps and endpoints. In particular, Both the claimed methods rely upon antibodies that bind B7-1 or B7-2. B7-1 and B7-2 differ in structure, expression and modes of action to such an extent and require non-coextensive searches to such an extent that they are patentably distinct.
- 6. Inventions I and II are different products. The inventions encompass antibodies that bind B7-1 and B7-
- 2. B7-1 and B7-2 differ in structure, expression and modes of action to such an extent and require non-coextensive searches to such an extent that they are patentably distinct.

Application/Control Number: 10/073,138

Art Unit: 1644

7. Because these inventions are distinct for the reasons given above and the search required for any group from Groups I-IV is not required for any other group from Groups I-IV and Groups I-IV have acquired a separate status in the art because the searches are not co-extensive and encompass divergent subject matter, restriction for examination purposes as indicated is proper.

- 8. This application contains claims directed to the following patentably distinct species of the claimed Invention III and IV: wherein the disease is:
 - A) an autoimmune disorder,
 - B) transplant rejection,
 - C) GVHD,
 - D) B cell lymphoma,
 - E) infectious diseases, or
 - F) inflammatory diseases.

These species are distinct because the pathological conditions differ in etiologies and therapeutic endpoints.

Applicant is required under 35 U.S.C. § 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 16 and 17 are generic for example.

In addition to electing Invention III or IV and a disease selected from (A) - (F), applicant is required to elect a specific species as it reads on (A) - (F) as disclosed on page 37, paragraph 1 of the instant specification.

9. Applicant is advised that a response to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 C.F.R. § 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. M.P.E.P. § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. § 103 of the other invention.

10. Applicant is advised that the response to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed.

Application/Control Number: 10/073,138

Art Unit: 1644

- 11. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 C.F.R. § 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a diligently-filed petition under 37 C.F.R. § 1.48(b) and by the fee required under 37 C.F.R. § 1.17(h).
- 12. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Phillip Gambel whose telephone number is (571) 272-0844. The examiner can normally be reached Monday through Thursday from 7:30 am to 6:00 pm. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Chan can be reached on (571) 272-0841.

The fax number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

PHUIP CAMBEL
Phillip Gambel, PhD.

Primary Examiner

Technology Center 1600

September 30, 2004